- ANSWER 1 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 1 L2
- AN 2004:490265 CAPLUS
- DN 141:52841
- Cloning and characterization of genes encoding culture filtrate antigens TT involved in protective immunity to M. tuberculosis, and use thereof as vaccines and in diagnosis
- Andersen, Peter; Skiot, Rikke; Oettinger, Thomas; Rasmussen, Peter Birk; IN Rosenkrands, Ida; Weldingh, Karin; Florio, Walter
- PA
- U.S. Pat. Appl. Publ., 109 pp., Cont.-in-part of U.S. 6,641,814. SO CODEN: USXXCO
- DTPatent
- English LA
- DANT CATTLE O

FAN.	CNT	9																	
	PAT	ENT	NO.			KIND		DATE			APPLICATION NO.						DATE		
							-									-			
ΡI	US	2004	1152	11		A1		2004		US 2003-620246					20030715				
	US	6641	814			B1		2003	1104		US	1998-	5073	9		1:	9980	330	
	ΕP	1449	922			A2		2004	0825		ΕP	2004-	7660	5		1	9980	401	
	ΕP	1449	922			A3		2004	1117										
		R:	AT,	BE,	CH,	DE,	DK	, ES,	FR,	GB,	GR	, IT,	LI,	LU,	NL,	SE,	MC,	PT,	
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	US .	JS 2002094336						2002	0718		US	2001-	7911	71		2	0010	220	
PRAI	DK	1997	-376			Α		1997	0402										
	US	1997	-4462	24P		P		1997	0418										
	DK	1997	-127	7		Α		1997	1110										
	US	1998	-7048	98P		P		1998	0105										
	US	1998	-5073	39		A2		1998	0330										
	DK	1998	-128	1		Α		1998	1008										
	EP	1998	-913	536		A3		1998	0401										

- The present invention is based on the identification and characterization AB of a number of M. tuberculosis derived antigens, isolated from culture filtrates of T cells from memory immune mice by T cell epitope mapping. The invention is directed to the polypeptides and immunol. active fragments thereof, the genes encoding them, immunol. compns. such as vaccines and skin test reagents containing the polypeptides. Another part of the invention is based on the surprising discovery that fusions between ESAT-6 and MPT59 are superior immunogens compared to each of the unfused proteins, resp. These antigens are suitable for use in vaccines and in diagnosis of infections.
- 1.2 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 2
- 2004:59568 CAPLUS ΔN
- 140:127185 DN
- Antigens from Mycobacterium as vaccine and uses in tuberculosis diagnosis TТ and treatment
- Andersen, Peter; Skjot, Rikke Louise Vinther; Okkels, Li Mei Meng; Brock, IN Inger; Oettinger, Thomas
- PA
- U.S. Pat. Appl. Publ., 27 pp., Cont.-in-part of U.S. Ser. No. 804,980. SO CODEN: USXXCO
- ידת Patent
- LΑ English

FAN.	CNT	9																		
	PAT	CENT 1	NO.			KIN)	DATE			APPLICATION NO.						DATE			
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ΡI	US	5 2004013685				A1 20040122				1	US 2001-872505						20010601			
EP 1449922						A2		20040825			EP 2004-76605					19980401				
	EP 1449922					A3		2004	0041117											
		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,		
			ΙE,	FI,	CY															
	WO 2001004151 WO 2001004151					A2		2001	0118	1	WO 2	O 2000-DK39				20000713				
						A3		2001	0712											
		W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,		
			CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,	HR,		
			HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KP,	KR,	KZ,	LC,	LK,	LR,	LS,	LT,		

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LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
             SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
             YU, ZA, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
             CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     US 2003147897
                         A1
                               20030807
                                           US 2001-804980
                                                                  20010313
PRAI DK 1997-1277
                         Α
                               19971110
     US 1998-70488P
                        P
                               19980105
                        B2
     US 1998-246191
                               19981230
     DK 1999-1020
                        Α
                               19990713
     US 1999-144011P
                        P
                               19990715
     US 2000-615947
                        A2
                               20000713
                        A2
     WO 2000-DK398
                               20000713
                        A2
     US 2001-804980
                               20010313
    US 1993-123182
     DK 1993-798
                        Α
                               19930702
                      B2
                               19930920
                        A2
                               19940701
     US 1995-465640
                        A1
                               19950605
     DK 1997-376
                        Α
                               19970402
     US 1997-44624P
                        Р
                               19970418
     EP 1998-913536
                        A3
                               19980401
     US 1999-289388
                        B2
                               19990412
AB
     The present invention is based on the identification and characterization
     of 3 antigens, including Rv2653c, Rv2654c and RD1-ORF5
     , from Mycobacterium tuberculosis. The invention is directed to the
     polypeptides and immunol. active fragments thereof, the genes encoding
     them, immunol. compns. such as diagnostic reagents containing the
     polypeptides. The invention related to diagnosing tuberculosis caused by
     virulent mycobacteria in an animal, including a human being. The
     invention related to treating tuberculosis using antigens isolated from
     Mycobacterium tuberculosis.
L_2
     ANSWER 3 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN
AN
     2004:60336 CAPLUS
DN
     140:144681
     Mycobacterium low oxygen-induced antigens and genes for vaccines or
ΤI
     diagnostics of tuberculosis
IN
     Andersen, Peter; Rosenkrands, Ida; Stryhn, Anette
PA
     Statens Serum Institut, Den.
SO
     PCT Int. Appl., 76 pp.
     CODEN: PIXXD2
DТ
     Patent
LA
     English
FAN.CNT 1
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                        KIND
                                          APPLICATION NO.
                               DATE
                                                                 DATE
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                               20040122
PΙ
     WO 2004006952
                        A2
                                           WO 2003-DK477
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     WO 2004006952
                        A3
                               20040318
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
            CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
            GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
            LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT,
            RO, RU, SD, SE, SG, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ,
            VN, YU, ZA, ZW
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
            KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
            FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
            BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
    EP 1523331
                              20050420 EP 2003-763613
                         A2
                                                                 20030708
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
    US 2004057963
                               20040325
                                          US 2003-617038
                        A1
                                                                  20030711
PRAI DK 2002-1098
                               20020713
                         Α
    US 2002-401725P
                         P
                               20020807
                         W
    WO 2003-DK477
                               20030708
AB
    The present invention is based on a number of M. tuberculosis derived
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proteins and protein fragments which are induced during the latent stage of infection characterized by low oxygen tension in the microenvironment of the infecting TB-bacteria. The invention is directed to the use of these polypeptides, immunol. active fragments thereof and the genes encoding them for immunol. compns. such as therapeutic vaccines and diagnostic reagents.

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L2
    ANSWER 4 OF 10 USPATFULL on STN
       2004:76186 USPATFULL
AN
      Therapeutic TB vaccine
TI
TN
      Andersen, Peter, Bronshoj, DENMARK
      Rosenkrands, Ida, Vaerlose, DENMARK
      Stryhn, Anette, Virum, DENMARK
PΙ
      US 2004057963
                        A1
                              20040325
      US 2003-617038
                        A1
                              20030711 (10)
AΙ
                         20020713
PRAI
      DK 2002-1098
      US 2002-401725P
                          20020807 (60)
DT
      Utility
FS
      APPLICATION
      HOWSON AND HOWSON, ONE SPRING HOUSE CORPORATION CENTER, BOX 457, 321
LREP
      NORRISTOWN ROAD, SPRING HOUSE, PA, 19477
CLMN
      Number of Claims: 22
ECL
      Exemplary Claim: 1
DRWN
      7 Drawing Page(s)
LN.CNT 6018
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      Therapeutic vaccines comprising polypeptides expressed during the latent
AB
       stage of mycobacteria infection are provided, as are multiphase
      vaccines, and methods for treating and preventing tuberculosis.
    ANSWER 5 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 3
1.2
    2003:696302 CAPLUS
ΑN
     139:229237
DN
     Protein and DNA sequences of antigens from Mycobacterium and uses in
ΤI
     tuberculosis diagnosis and treatment
    Andersen, Peter; Weldingh, Karin; Hansen, Christina Veggerby; Florio,
IN
    Walter; Okkels, Li Mei Meng; Skjot, Rikke Louise Vinther; Rasmussen, Peter
    Rirk
    Den.
PA
    U.S. Pat. Appl. Publ., 53 pp., Cont.-in-part of U.S. Ser. No. 60,428.
SO
     CODEN: USXXCO
DT
     Patent
LA
    English
FAN.CNT 10
                                           APPLICATION NO.
                                                                 DATE
    PATENT NO.
                       KIND
                               DATE
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                               20030904
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PΙ
    US 2003165525
                        A1
    US 6641814
                        B1
                               20031104
                                           US 1998-50739
                                                                 19980330
     EP 1449922
                                           EP 2004-76605
                                                                 19980401
                        A2
                               20040825
     EP 1449922
                         A3
                               20041117
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI, CY
     US 2002094336
                                           US 2001-791171
                                                                  20010220
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                               20020718
                        Α
PRAI DK 1997-376
                               19970402
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     US 1997-44624P
                               19970418
                        Α
     DK 1997-1277
                               19971110
                        P
     US 1998-70488P
                               19980105
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     US 1998-50739
                               19980330
                        Α
     DK 1998-1281
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     US 2001-791171
                         B2
                               20010220
     US 2002-60428
                         A2
                               20020129
     EP 1998-913536
                         A3
                               19980401
     The present invention is based on the identification and characterization
AB
     of 9 antigens, including Rv0652/CFP16, Rv2462c/TB51, Rv1984c/CFP21,
     Rv2185c/TB16, Rv1636/TB15A, Rv3451/CFP23, Rv3872/RD1-ORF3, Rv3354/CFP8A
     and Rv2623/TB32, from Mycobacterium tuberculosis. The invention is
     directed to the polypeptides and immunol. active fragments thereof, the
     genes encoding them, immunol. compns. such as diagnostic reagents containing
     the polypeptides. The invention related to diagnosing tuberculosis caused
     by virulent mycobacteria, e.g. by Mycobacterium tuberculosis,
     Mycobacterium africanum or Mycobacterium bovis, in an animal, including a
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human being. The invention related to treating tuberculosis using antigens isolated from Mycobacterium tuberculosis.

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ANSWER 6 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 4
L2
AN
     2003:609858 CAPLUS
DN
     139:163576
     Mycobacterium tuberculosis antigens for diagnosis, prevention and
     treatment of infections caused by species of the tuberculosis complex
IN
     Andersen, Peter; Skjot, Rikke Louise Vinther
PA
     U.S. Pat. Appl. Publ., 135 pp., Cont.-in-part of U.S. Ser. No. 289,388,
SO
     abandoned.
     CODEN: USXXCO
DT
     Patent
LΑ
     English
FAN.CNT 9
     PATENT NO.
                       KIND
                               DATE
                                         APPLICATION NO.
                                                                DATE
                                          -----
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PΙ
     US 2003147897
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                                          US 2001-804980
                                                                20010313
     WO 9501441
                        A1
                               19950112
                                          WO 1994-DK273
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            NO, NZ, PL, PT, RO, RU, SD, SE, SI, SK, TJ, TT, UA
        RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE,
            BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG
                                          EP 2004-77505
     EP 1508339
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    US 5955077
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                               19990921
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     EP 1449922
                         A2
                               20040825
                                          EP 2004-76605
                                                                 19980401
     EP 1449922
                         A3
                               20041117
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, FI, CY
     US 2004013685
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                               20040122
                                          US 2001-872505
                                                                 20010601
PRAI DK 1993-798
    DK 1993-750
US 1993-123182
                        Α
                               19930702
                       B2
                               19930920
     WO 1994-DK273
                       A2
                               19940701
    US 1995-465640
                        A1
                               19950605
    DK 1997-376
                        Α
                               19970402
    US 1997-44624P
                        P
                               19970418
                       A 19971110
    DK 1997-1277
    US 1998-70488P
                        P
                               19980105
    US 1999-289388
                       B2
                               19990412
    EP 1994-919574
                        A3
                               19940701
    EP 1998-913536
                        A3
                               19980401
    US 1998-246191
                        B2
                               19981230
    DK 1999-1020
                        Α
                               19990713
    US 1999-144011P
                        P
                               19990715
    US 2000-615947
                        A2
                               20000713
     WO 2000-DK398
                        A2
                               20000713
    US 2001-804980
                        A2
                               20010313
AB
     The present invention is based on the identification and characterization
     of a number of novel M. tuberculosis derived proteins and protein fragments,
     e.g. TB10.3 (ORF7-1 or Rv3019c), TB10.4 (CFP7 or Rv0288) and TB12.9
     (ORF7-2 or Rv3017c), ESAT-6, MPT64, CFP10, RD1-ORF5,
     RD1-ORF2, Rv1036, Ag85A, Ag85B, Ag85C, 19 kDa lipoprotein, MPT32, MPB59
     and \alpha-crystallin. The invention is directed to the polypeptides and
     immunol. active fragments thereof, the genes encoding them, immunol.
     compns. such as vaccines and skin test reagents containing the polypeptides.
    ANSWER 7 OF 10 USPATFULL on STN
L2
AN
      2003:291011 USPATFULL
TI
      Nucleic acids fragments and polypeptide fragments derived from M.
      tuberculosis
      Andersen, Peter, Br.o slashed.nsh.o slashed.j, DENMARK
IN
      Nielsen, Rikke, Frederiksberg, DENMARK
      Oettinger, Thomas, Hellerup, DENMARK
      Rasmussen, Peter Birk, K.o slashed.benhaven, DENMARK
      Rosenkrands, Ida, K.o slashed.benhaven, DENMARK
```

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Weldingh, Karin, K.o slashed.benhaven, DENMARK
       Florio, Walter, Frederiksberg, DENMARK
PΑ
       Statens Serum Institut, Copenhagen, DENMARK (non-U.S. corporation)
PΙ
                               20031104
       US 6641814
                          В1
ΑI
       US 1998-50739
                               19980330 (9)
PRAI
       DK 1997-376
                           19970402
       US 1997-44624P
                           19970418 (60)
DΤ
       Utility
FS
       GRANTED
      Primary Examiner: Swartz, Rodney P
EXNAM
       Frommer Lawrence & Haug, Kowalski, Thomas J.
LREP
       Number of Claims: 43
CLMN
ECL
       Exemplary Claim: 1
       6 Drawing Figure(s); 6 Drawing Page(s)
DRWN
LN.CNT 5870
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention is based on the identification and
       characterization of a number of M. tuberculosis derived novel proteins
       and protein fragments (SEQ ID NOs: 2, 4, 6, 8, 10, 12, 14, 16, 17-23,
       42, 48, 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72-86, 88, 90, 92,
       94, 141, 143, 145, 147, 149, 151, 153, and 168-171). The invention is
       directed to the polypeptides and immunologically active fragments
       thereof, the genes encoding them, immunological compositions such as
       vaccines and skin test reagents containing the polypeptides. Another
       part of the invention is based on the surprising discovery that fusions
       between ESAT-6 and MPT59 are superior immunogens compared to each of the
       unfused proteins, respectively.
L2
     ANSWER 8 OF 10 USPATFULL on STN
       2002:314395 USPATFULL
AN
ΤI
       Hybrids of M. tuberculosis antigens
       Andersen, Peter, Bronshoj, DENMARK
IN
       Olsen, Anja Weinreich, Soborg, DENMARK
       Skjot, Rikke Louise Vinther, Hedehusene, DENMARK
       Rasmussen, Peter Birk, Frederiksberg, DENMARK
PΙ
       US 2002176867
                          A1
                               20021128
       US 2001-805427
                          A1
                               20010313 (9)
ΑI
       Continuation-in-part of Ser. No. US 1998-246191, filed on 30 Dec 1998,
RLI
       ABANDONED
                           19971110
PRAI
       DK 1997-1277
       US 1998-70488P 19980105 (60)
                           19970418 (60)
       US 1997-44624P
DT
       Utility
       APPLICATION
FS
       Thomas J. Kowalski, c/o FROMMER LAWRENCE & HAUG LLP, 745 Fifth Avenue,
LREP
       New York, NY, 10151
CLMN
       Number of Claims: 25
ECL
       Exemplary Claim: 1
DRWN
       10 Drawing Page(s)
LN.CNT 2157
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention discloses fusion proteins of the immunodominant
AB
       antigens ESAT-6 and Ag85B from Mycobacterium tuberculosis or homologues
       thereof, and a tuberculosis vaccine based on the fusion proteins, which
       vaccine induces efficient immunological memory.
     ANSWER 9 OF 10 USPATFULL on STN
L2
       2002:178550 USPATFULL
AN
       Nucleic acid fragments and polypeptide fragments derived from M.
ΤI
       tuberculosis
IN
       Andersen, Peter, Bronshoj, DENMARK
       Nielsen, Rikke, Frederiksberg C, DENMARK
       Oettinger, Thomas, Hellerup, DENMARK
       Rasmussen, Peter Birk, Kobenhaven O, DENMARK
       Rosenkrands, Ida, Kobenhaven O, DENMARK
       Weldingh, Karin, Kobenhaven N, DENMARK
       Florio, Walter, Frederiksberg C, DENMARK
PA
       STATENS SERUM INSTITUT (non-U.S. corporation)
ΡI
       US 2002094336
                          A1
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ΑI
       US 2001-791171
                         A1
                               20010220 (9)
RLI
       Division of Ser. No. US 1998-50739, filed on 30 Mar 1998, PENDING
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       DK 1997-376
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       DK 1997-1277
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       US 1997-44624P
                           19970418 (60)
       US 1998-70488P
                          19980105 (60)
DT
       Utility
FS
       APPLICATION
LREP
       FROMMER LAWRENCE & HAUG LLP, 745 FIFTH AVENUE, NEW YORK, NY, 10151
CLMN
       Number of Claims: 53
       Exemplary Claim: 1
ECL
DRWN
       6 Drawing Page(s)
LN.CNT 6134
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention is based on the identification and
AB
       characterization of a number of M. tuberculosis derived novel proteins
       and protein fragments (SEQ ID NOs: 2, 4, 6, 8, 10, 12, 14, 16, 17-23,
       42, 48, 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72-86, 88, 90, 92,
       94, 141, 143, 145, 147, 149, 151, 153, and 168-171). The invention is
       directed to the polypeptides and immunologically active fragments
       thereof, the genes encoding them, immunological compositions such as
       vaccines and skin test reagents containing the polypeptides. Another
       part of the invention is based on the surprising discovery that fusions
       between ESAT-6 and MPT59 are superior immunogens compared to each of the
       unfused proteins, respectively.
     ANSWER 10 OF 10 CAPLUS COPYRIGHT 2006 ACS on STN
L2
     1998:684968 CAPLUS
AN
DN
     129:300060
     Novel antigens of Mycobacterium tuberculosis culture filtrates and the
TI
     genes encoding and their diagnostic and prophylactic use
IN
     Andersen, Peter; Nielsen, Rikke; Rosenkrands, Ida; Weldingh, Karin;
     Rasmussen, Peter Birk; Oettinger, Thomas; Florio, Walter
PΑ
     Statens Serum Institut, Den.
SO
     PCT Int. Appl., 264 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 9
     PATENT NO.
                         KIND
                               DATE
                                           APPLICATION NO.
                                                                  DATE
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                                                                  19980401
                                19981008
                                            WO 1998-DK132
ΡI
                         A1
     WO 9844119
         W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
             DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG,
             KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
             NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
             UA, UG, US, UZ, VN, YU, ZW
         RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES,
             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI,
             CM, GA, GN, ML, MR, NE, SN, TD, TG
                                            CA 1998-2285625
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     AU 9868204
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     EP 972045
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             IE, FI
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                                            EP 2004-76605
                                                                   19980401
                         A2
                                20040825
     EP 1449922
                         A3
                                20041117
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             IE, FI, CY
     CA 2319380
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AB Culture filtrate antigens of Mycobacterium tuberculosis are characterized and cDNAs encoding them are cloned. Some of the proteins are antigenic and suitable for use in vaccines and in diagnosis of infections, e.g. skin tests. A fusion protein of two of these antigens is a superior immunogen compared to the unfused proteins. Individual antigens from culture filtrates were identified by T cell mapping using T cells from memory immune mice. Genes for individual antigens were then cloned by screening a \(\lambda\gamma\)11 expression vector with monoclonal antibodies. Manufacture of individual antigens with hexahistidine affinity labels is described.

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